The listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

A compound of formula I 1. (Currently Amended)

$$R^{1}$$
 R^{2}
 R^{3}

in which

 R^1 denotes (CH₂), Het1, or (CH₂), Ar, or eyeloalkyl having 3 to 7 C atoms,

is 4-pyridyl, thiophen-2-yl or thiophen-3-yl, which is unsubstituted or mono-Het1 or polysubstituted by CN, A and/or Hal,

 \mathbb{R}^2 denotes Het2 (CH₂)_nHet2, (CH₂)_nAr, or cycloalkyl having 3 to 7 C atoms,

is 2- or 3-furanyl, which is unsubstituted or mono- or polysubstituted by A Het2 and/or Hal,

one of the radicals R³ or R⁴ denotes H, and the other of the radicals R³ or R⁴ R^3, R^4 denotes denote-H, (CH₂)_nCO₂R⁵, (CH₂)_nCOHet3, CHO, (CH₂)_nOR⁵, $(CH_2)_nHet3$, $(CH_2)_nN(R^5)_2$, CH=N-OA, $CH_2CH=N-OA$, $(CH_2)_nNHOA$,

 $(CH_2)_nN(R^5)Het3$, $(CH_2)_nCH=N-Het3$, $(CH_2)_nOCOR^5$,

(CH₂)_nN(R⁵)CH₂CH₂OR⁵, <math>(CH₂)_nN(R⁵)CH₂CH₂OCF₃,

 $(CH_2)_nN(R^5)C(R^5)HCOOR^5$, $(CH_2)_nN(R^5)CH_2COHet_2$, $(CH_2)_nN(R^5)CH_2$

 $Het\underline{3}, (CH_2)_nN(R^5)CH_2CH_2 \ Het\underline{3}, (CH_2)_nN(R^5)CH_2CH_2N(R^5)CH_2COOR^5,$

 $(CH_2)_nN(R^5)CH_2CH_2N(R^5)_2$, $CH=CHCOOR^5$, $CH=CHCH_2NR^5$ Het3,

CH=CHCH₂N(R⁵)₂, CH=CHCH₂OR⁵ or (CH₂)_nN(R⁵)Ar,

with the proviso that in each case one of the radicals R³ or R⁴ denotes H,

is 1-piperidyl, 1-piperazyl, 1-(4-methyl)piperazyl, 4-methylpiperazin-1-Het3

ylamine, 1-pyrrolidinyl, 1-pyrazolidinyl, 1-(2-methyl)pyrazolidinyl, 1-imidazo-

lidinyl or 1-(3-methyl)imidazolidinyl or 4-pyridyl, which may be unsubstituted

or substituted by one or more CN groups, 2- or 4-pyridazyl, 2-, 4- or

5-pyrimidyl, or 2- or 3-pyrazinyl or one of the following groups

R⁵ denotes H or A,

A denotes straight-chain or branched alkyl or alkoxy having 1 to 10 C atoms, alkenyl or alkoxyalkyl having 2 to 10 C atoms,

Het denotes a saturated, unsaturated or aromatic mono- or bicyclic heterocyclic or linear or branched organic radical containing one or more heteroatoms which is unsubstituted or mono- or polysubstituted by A and/or Hal,

Ar denotes a phenyl radical which is unsubstituted or mono- or polysubstituted by A and/or Hal, OR⁵, OOCR⁵, COOR⁵, CON(R⁵)₂, CN, NO₂, NH₂, NHCOR⁵, CF₃ or SO₂CH₃,

n denotes 0, 1, 2, 3, 4 or 5,

Hal denotes F, Cl, Br or I, and

X denotes N, or
in the case where R¹ denotes one of the following groups

in which R denotes H or an alkyl group having 1 to 6 C atoms, and/or R² denotes

in which R denotes H or an alkyl group having 1 to 6 C atoms, alternatively denotes CH,

or an enantiomer, racemate, or a mixture of enantiomers thereof, or a pharmaceutically acceptable salt thereof.

2. (Previously Presented) A compound of formula I according to Claim 1, in which R¹ denotes phenyl, 2-, 3- or 4-cyanophenyl, 2-, 3- or 4-fluorophenyl, 2-, 3- or 4-methyl-, -ethyl-, -n-propyl- or -n-butylphenyl, 2,3-, 2,4-, 2,5-, 2,6-, 3,4-, 3,5- or 3,6-difluoro-, -dichloro- or -dicyanophenyl, 3,4,5trifluorophenyl, 3,4,5-trimethoxy- or -triethoxyphenyl, thiophen-2-yl or thiophen-3-yl.

3. (Previously Presented) claim 1, in which R³ denotes H.

A compound of formula I according to

4. (Previously Presented) claim 1, in which R⁴ denotes H.

A compound of formula I according to

- 5. (Cancelled)
- 6. (Previously Presented) claim 1, in which X denotes N.

A compound of formula I according to

7. (Currently Amended) is of formula IA, IB, IC, ID, IE or IF

A compound according to claim 1, which

$$R^{2}$$
 R^{2}
 R^{2}
 R^{2}
 R^{3}
 R^{4}
 R^{2}
 R^{4}
 R^{2}
 R^{4}
 R^{2}
 R^{4}
 R^{4}
 R^{4}
 R^{5}
 R^{2}
 R^{4}
 R^{5}
 R^{5}

in which

R¹, R², X and A are as defined for the compound of formula I.

(CH₂)_nAr, or cycloalkyl having 3 to 7 C atoms,

denotes (CH₂)_nHet, (CH₂)_nAr, or cycloalkyl having 3 to 7 C atoms,

A denotes straight chain or branched alkyl or alkoxy having 1 to 10 C atoms,

alkenyl or alkoxyalkyl having 2 to 10 C atoms,

Het denotes a saturated, unsaturated or aromatic mono- or bicyclic heterocyclic or linear or branched organic radical containing one or more heteroatoms which is unsubstituted or mono- or polysubstituted by A and/or Hal,

Ar denotes a phenyl radical which is unsubstituted or mono- or

polysubstituted by A and/or Hal, OR^5 , $OOCR^5$, $COOR^5$, $CON(R^5)_2$, CN, NO_2 , NH_2 , $NHCOR^5$, CF_3 or SO_2CH_3 ,

R⁵ denotes H or A,

n denotes 0, 1, 2, 3, 4 or 5,

Hal denotes F, Cl, Br or I, and

X denotes N, or

in the case where R¹-denotes

in which R denotes H or an alkyl group having 1 to 6 C atoms, and/or R²-denotes

in which R denotes H or an alkyl group having 1 to 6 C atoms, alternatively denotes CH,

or a salt thereof.

8. (Previously Presented) A process for preparing a compound of formula IA according to claim 7

comprising reacting a compound of formula II

$$R^1$$
 NHNH₂

or an acid-addition salt thereof, in which R¹ and X have the meanings indicated for the compound of formula IA, with a compound of formula III

$$\mathbb{R}^2$$
 \mathbb{N} \mathbb{N}

in which

A and R² have the meanings indicated for the compound of formula IA, and/or

a basic compound of formula IA is converted into one of its salts by treatment with an acid.

9. (Previously Presented) A process for preparing a compound of

formula IB according to claim 7

in which R¹, R², R³, R⁴, X and A have the meanings indicated for the compound of formula IB,

comprising reacting a compound of formula II

or an acid-addition salt thereof, in which $R^1 \ and \ X \ have the meanings indicated for the compound of formula IB,$ with a compound of formula IV

$$\mathbb{R}^2$$
 \mathbb{I}^2

in which

A and R² have the meanings indicated for the compound of formula IB, and/or

a basic compound of formula IB is converted into one of its salts by treatment with an acid.

- 10. (Previously Presented) A pharmaceutical composition comprising a compound of formula I according to claim 1 and a pharmaceutically acceptable carrier.
- 11. (Previously Presented) A method for the treatment of a disease which can be influenced by the binding of a compound of formula I to 5 HT receptors, comprising administering to a subject in need thereof an effective amount of a pharmaceutical

composition according to claim 10.

- 12. (Previously Presented) A method for antagonizing a 5-HT receptor, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 10.
- 13. (Previously Presented) A method for antagonizing a 5-HT_{2A} receptor, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 10.
 - 14. (Cancelled)
- 15. (Previously Presented) A process for preparing a pharmaceutical composition according to claim 10, comprising mixing together a compound of formula I and a pharmaceutically acceptable carrier.
- 16. (Currently Amended) A method for the treatment of psychoses, a neurological disorder, amyotrophic lateral sclerosis, eating disorder, bulimia, anorexia nervosa, premenstrual syndrome and/or for positively influencing obsessive compulsive disorder, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 10.

17-22. (Cancelled)

23. (Currently Amended) A compound of claim 1, in which

R¹ denotes Het1 or Ar,

R² denotes Het or Ar,

R³, R⁴ denote H, (CH₂)_nCO₂R⁵, CH=N-OA, CH₂CH=N-OA, (CH₂)_nNHOA,

(CH₂)_nN(R⁵)Het, (CH₂)_nCH=N-Het, (CH₂)_nOCOR⁵, (CH₂)_nN(R⁵)CH₂CH₂OR⁵,

(CH₂)_nN(R⁵)CH₂CH₂OCF₃, (CH₂)_nN(R⁵)C(R⁵)HCOOR⁵;

(CH₂)_nN(R⁵)CH₂COHet, (CH₂)_nN(R⁵)CH₂Het, (CH₂)_nN(R⁵)CH₂CH₂Het,

(CH₂)_nN(R⁵)CH₂COHet, (CH₂)_nN(R⁵)CH₂Het, (CH₂)_nN(R⁵)CH₂CH₂N(R⁵)₂;

CH=CHCOOR⁵, CH=CHCH₂NR⁵Het, CH=CHCH₂N(R⁵)₂, CH=CHCH₂OR⁵ or

(CH₂)_nN(R⁵)Ar, with the proviso that in each case one of the radicals R³ or R⁴ denotes H,

R⁵ denotes H or A,

A denotes straight chain or branched-alkyl or alkoxy having 1 to 10 C atoms, alkenyl or alkoxyalkyl having 2 to 10 C atoms,

Het denotes a saturated, unsaturated or aromatic mono- or bicyclic heterocyclic or linear or branched organic radical containing one or more heteroatoms which is unsubstituted or mono- or polysubstituted by A and/or Hal,

Ar denotes a phonyl radical which is unsubstituted or mono or polysubstituted by A and/or Hal, OR⁵, OOCR⁵, COOR⁵, CON(R⁵)₂, CN, NO₂, NH₂, NHCOR⁵, CF₃ or SO₂CH₃,

n denotes 0, 1, 2 or 3,

Hal denotes F, Cl, Br or I, and

X denotes N, or

in the case where R⁴ denotes

in which R denotes H or an alkyl group having 1 to 6 C atoms, and/or R² denotes

in which R denotes H or an alkyl group having 1 to 6 C atoms, alternatively denotes CH.

24-27. (Cancelled)

- 28. (Previously Presented) A method for antagonizing a 5-HT_{2A} receptor in vitro, comprising administering to said 5-HT_{2A} receptor an effective amount of a compound according to claim 1.
- 29. (Currently Amended) A method for the treatment of psychoses, amyotrophic lateral sclerosis, bulimia, anorexia nervosa, premenstrual syndrome and/or for positively influencing obsessive compulsive disorder, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 10.
- 30. (New) A method for the treatment of amyotrophic lateral sclerosis, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 10.
- 31. (New) A method for the treatment of bulimia, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 10.
 - 32. (New) A method for the treatment of anorexia nervosa, comprising

administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 10.

- 33. (New) A method for the treatment of premenstrual syndrome, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 10.
- 34. (New) A method for positively influencing obsessive compulsive disorder, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 10.
- 35. (New) A compound of formula I according to Claim 1, in which one of the radicals R^3 or R^4 denotes H and the other of the radicals R^3 or R^4 denotes $(CH_2)_nCO_2R^5$, $(CH_2)_nCOHet3$, $(CH_2)_nHet3$, $(CH_2)_nN(R^5)Het3$, $(CH_2)_nCH=N-Het3$, $(CH_2)_nN(R^5)CH_2CH_2OR^5$, $(CH_2)_nN(R^5)CH_2CH_2OCF_3$, $(CH_2)_nN(R^5)C(R^5)HCOOR^5$, $(CH_2)_nN(R^5)CH_2COHet3$, $(CH_2)_nN(R^5)CH_2COOR^5$, $(CH_2)_nN(R^5)CH_2COOR^5$, $(CH_2)_nN(R^5)COHet3$, or $(COHe_2)_nN(R^5)COOR^5$, wherein n is 1, 2, 3, 4 or 5.
- 36. (New) A compound of formula I according to Claim 1, in which one of the radicals R^3 or R^4 denotes H and the other of the radicals R^3 or R^4 denotes (CH₂)_nHet3, wherein n is 1, 2, 3, 4 or 5.